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NEWS 18 Aug 08 NTIS has been reloaded and enhanced
                 Aquatic Toxicity Information Retrieval (AQUIRE)
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                 now available on STN
                 IFIPAT, IFICDB, and IFIUDB have been reloaded
NEWS 20 Aug 19
                 The MEDLINE file segment of TOXCENTER has been reloaded
         Aug 19
NEWS 21
                 Sequence searching in REGISTRY enhanced
NEWS 22
         Aug 26
                 JAPIO has been reloaded and enhanced
NEWS 23
         Sep 03
                 Experimental properties added to the REGISTRY file
NEWS 24
         Sep 16
                 Indexing added to some pre-1967 records in CA/CAPLUS
NEWS 25
         Sep 16
                 CA Section Thesaurus available in CAPLUS and CA
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         Sep 16
                 CASREACT Enriched with Reactions from 1907 to 1985
NEWS 27
         Oct 01
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=> s "Apo-I" L1 695 "APO-I"

=> s 11 and conjugate L2 3 L1 AND CONJUGATE

=> dup reomve 12
ENTER REMOVE, IDENTIFY, ONLY, OR (?):remove
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PROCESSING COMPLETED FOR L2
L3 3 DUP REMOVE L2 (0 DUPLICATES REMOVED)

=> d 13 1-3 cbib abs

L3 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2002 ACS
1989:111335 Document No. 110:111335 Reagents for reactivation by cofactor of
an apoenzyme-antigen conjugate in an enzyme immunoassay method.
Siddiqi, Iqbal; Mangan, Ciaron (Battelle Memorial Institute, Switz.).
Eur. Pat. Appl. EP 274343 A1 19880713, 9 pp. DESIGNATED STATES: R: AT,
BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE. (English). CODEN:
EPXXDW. APPLICATION: EP 1987-810005 19870106.

AB A homogeneous EIA is described in which an apoenzyme conjugated to an antigen is regenerated by addn. of a cofactor, and regeneration of the holoenzyme is inhibited by complexing the antigen to an antibody. Alk. phosphatase (I), conjugated to a rabbit antibody to human IgG, was completely inhibited by 10-2M 1,10-phenanthroline. The activity was 10-20% restored by addn. of 0.3M Zn2+, as compared to 100% for unconjugated I; thus conjugation altered the kinetics of reactivation, but reactivation was sufficient for use of the conjugate in an EIA.

When the apo-I-antibody conjugate was incubated with donkey anti-rabbit IgG antiserum, the greater the amt. of antiserum in the assay medium, the less the I activity was rescued by Zn2+. Addn. of competing unlabeled rabbit IgG (analyte) allowed restoration of I activity by Zn2+.

- L3 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2002 ACS
- 1985:518335 Document No. 103:118335 Purification and characterization of cytosolic liver protein facilitating heme transport into apocytochrome b5 from mitochondria. Evidence for identifying the heme transfer protein as belonging to a group of glutathione S-transferases. Senjo, Masanori; Ishibashi, Teruo; Imai, Yoh (Sch. Med., Hokkaido Univ., Sapporo, 060, Japan). J. Biol. Chem., 260(16), 9191-6 (English) 1985. CODEN: JBCHA3. ISSN: 0021-9258.
- The transport of protoheme from mitochondria to apocytochrome b5 (AB apo-I) was examd. by incubation of fresh rat liver mitochondria with apo-I and cytosol. The heme-transfer protein was purified .apprx.133- to 140-fold from rat liver cytosol, with a 43% yield by procedures which included Sephadex G-75 and CM-cellulose column chromatog. The final prepn. was apparently homogeneous by SDS-polyacrylamide gel electrophoresis. Its native form was a dimeric protein with a mol. wt. of 45,000 which consisted of a subunit with a mol. wt. of 23,000. Heme transfer depended on the concn. of mitochondria (donor), apo-I (acceptor), and purified transfer protein. Omission of any of these components led to an almost complete loss of the transfer activity. The transport of mitochondrial protoheme was rapid and approx. linear for .ltoreq.1.5 min, after which it became satd. When the functional capacity was tested by the NADH-cytochrome c reductase system, the reconstituted I expressed its complete original catalytic properties, as well as its characteristic absorption spectra for the hemoprotein. Furthermore, the detailed physicochem. and immunol. characterization of the transfer protein indicated that the protein was identical with sol. glutathione S-transferase (II), which conjugates glutathione with a variety of electrophilic compds. At least 1 of the II isoenzymes obsd. was II-C2 by immunopptn. with various anti-II isoenzyme antibodies.
- L3 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2002 ACS
- 1981:202859 Document No. 94:202859 Inhibition of kynureninase (L-kynurenine hydrolase, EC 3.7.1.3) by estrone sulfate: an alternative explanation for abnormal results of tryptophan load tests in women receiving estrogenic steroids. Bender, David A.; Wynick, David (Courtauld Inst. Biochem., Middlesex Hosp. Med. Sch., London, W1P 7PN, Engl.). Br. J. Nutr., 45(2), 269-75 (English) 1981. CODEN: BJNUAV. ISSN: 0007-1145.

 AB Rat liver kynureninase (EC 3.7.1.3) (I) [9024-78-6], in a partially
- AB Rat liver kynureninase (EC 3.7.1.3) (I) [9024-78-6], in a partially purified cofactor-free prepn., was uncompetitively inhibited by estrone sulfate [481-97-0] with respect to pyridoxal phosphate and competitively inhibited with respect to kynurenine with a Ki of 82 .mu.M. Addn. of a satg. concn. of pyridoxal phosphate to unfractionated liver homogenates increased I activity .apprx.5-fold, indicating the presence of a relatively large amt. of apo-I. It was suggested that the abnormal results of tryptophan load tests in women receiving estrogens may be the result of I inhibition by estrogen conjugates and that estrogen-induced vitamin B6 [8059-24-3] deficiency probably does not occur.
- => s l1 and ortholog L4 0 L1 AND ORTHOLOG
- => s human apolipoprotein AI L5 300 HUMAN APOLIPOPROTEIN AI

=> s 15 and cynomolgus monkey L6 0 L5 AND CYNOMOLGUS MONKEY

=> s 15 and macaca fascicularis L7 0 L5 AND MACACA FASCICULARIS

=> s 15 and PEG modified

L8 0 L5 AND PEG MODIFIED

=> s 15 and polythylene glycol L9 0 L5 AND POLYTHYLENE GLYCOL

=> dup remove 15
PROCESSING COMPLETED FOR L5
L10 150 DUP REMOVE L5 (150 DUPLICATES REMOVED)

=> s 110 and conjugate L11 1 L10 AND CONJUGATE

=> d l11 cbib abs

L11 ANSWER 1 OF 1 EMBASE COPYRIGHT 2002 ELSEVIER SCI. B.V.
92269493 EMBASE Document No.: 1992269493. Rapid screening method for
polymorphism of group A apolipoproteins. Harake B.; Caines P.S.M.; Thibert
R.J.. Department of Chemistry/Biochemistry, University of Windsor, Windsor,
Ont. N9B 3P4, Canada. Journal of Clinical Laboratory Analysis 6/5
(290-296) 1992.

ISSN: 0887-8013 CODEN: JCANEM Pub Country: United States Language:

ISSN: 0887-8013. CODEN: JCANEM. Pub. Country: United States. Language: English. Summary Language: English.

AΒ Polymorphism of apolipoproteins AI and AII (apo AI and apo AII) can be easily investigated in plasma by a simple method involving a 30-min incubation of EDTA plasma in the presence of urea, dithiothreitol, and Nonidet P-40 followed by subsequent isoelectric focusing (IEF). The sample (2 .mu.L) was applied to an ultrathin flat acrylamide gel of pH range 4-6, and focused using a Bio-Rad.RTM. Mini IEF Cell for 1.5 h at a maximum of 500 V. Coomassie Blue R-250 was used to visualize the apolipoproteins. To verify the identity of the different apolipoproteins after IEF, the gel was immunofixed directly with anti-3apo AI, or immunoblotted on polyvinylidene difluoride (PVDF) membrane using monospecific antibodies to apo AI and apo AII and an anti-immunoglobulin-alkaline phosphatase conjugate. High-density lipoprotein (HDL) was used as a standard for Apo AI variants. Employing these techniques, human plasma apo AI was resolved into one major band (apo AIO, pI 5.54), and four minor bands identified as apo AI+2 (pI 5.75), apo AI+1 (pI 5.66), apo AI-1 (pI 5.45), and apo AI-2 (pI 5.34). Apo AII was resolved into one major isoprotein designated as apo AII0 (pI 4.87), and two minor isoforms apo AII+1 and apo AII-1 which focused at pIs of 5.18 and 4.58, respectively. The results showed that these methods can be used to identify apo AI and AII isoforms without prior ultracentrifugation to isolate the HDL. The entire procedure, including IEF, fixation (chemical or immunofixation), and staining, can be accomplished in 5 h compared to 2 days using previously reported technique. The identification and characterization of human apolipoprotein AI and AII isoforms is

important in clinical practice, e.g., diagnosis of tangier disease, and may be useful in studying structure—function relationships of these apoproteins.

=> s 110 and copolymer L12 0 L10 AND COPOLYMER (FILE 'HOME' ENTERED AT 19:04:28 ON 07 OCT 2002)

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FILE 'MEDLINE, EMBASE, BIOSIS, SCISEARCH, CAPLUS' ENTERED AT 19:04:43 ON
     07 OCT 2002
             695 S "APO-I"
1.1
              3 S L1 AND CONJUGATE
L2
               3 DUP REMOVE L2 (0 DUPLICATES REMOVED)
L3
              0 S L1 AND ORTHOLOG
L4
            300 S HUMAN APOLIPOPROTEIN AI
L5
              0 S L5 AND CYNOMOLGUS MONKEY
L6
              0 S L5 AND MACACA FASCICULARIS
L7
              0 S L5 AND PEG MODIFIED
L8
              0 S L5 AND POLYTHYLENE GLYCOL
L9
            150 DUP REMOVE L5 (150 DUPLICATES REMOVED)
L10
              1 S L10 AND CONJUGATE
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              0 S L10 AND COPOLYMER
L12
=> s 11 and macaca fascicularis
             0 L1 AND MACACA FASCICULARIS
=> s polymer
      1510697 POLYMER
=> s 114 and water soluble
L15
         37443 L14 AND WATER SOLUBLE
=> s 115 adn polyethylene glycol
MISSING OPERATOR L15 ADN
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nested terms that are not separated by a logical operator.
=> s 115 and polyethylene glycol
          2874 L15 AND POLYETHYLENE GLYCOL
=> s 116 and monomethyoxy polythylene glycol
             0 L16 AND MONOMETHYOXY POLYTHYLENE GLYCOL
L17
=> s 116 adn dextran
MISSING OPERATOR L16 ADN
The search profile that was entered contains terms or
nested terms that are not separated by a logical operator.
=> s 116 and dextran
L18
          228 L16 AND DEXTRAN
=> s 118 and ApoAI
            0 L18 AND APOAI
L19
=> s 118 and peptide
            35 L18 AND PEPTIDE
=> dup remove 120
PROCESSING COMPLETED FOR L20
             35 DUP REMOVE L20 (0 DUPLICATES REMOVED)
=> d 121 1-35 cbib abs
L21 ANSWER 1 OF 35 CAPLUS COPYRIGHT 2002 ACS
2002:185399 Document No. 136:229029 Method for precipitating mono and
     multiple layers of organophosphoric and organophosphonic acids and the
     salts thereof in addition to use thereof. Hofer, Rolf; Pawlak, Michael; Textor, Marcus; Schuermann-Mader, Eveline; Ehrat, Markus; Tosatti, Samuele
     (Zeptosens A.-G., Switz.). PCT Int. Appl. WO 2002020873 A2 20020314, 88
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pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (German). CODEN: PIXXD2. APPLICATION: WO 2001-EP10077 20010831. PRIORITY: CH 2000-1732 20000905.

The invention relates to a method for pptg. mono or multiple layers of organophosphoric acids of general formula (I(A)) Y-B-OPO3 H2 (IA) or organophosphonic acids of general formula (I(B)) Y-B-PO3 H2 (IB) and the salts thereof, wherein B is an alkyl, alkenyl, alkynyl, aryl, aralkyl, hetaryl or hetaryl alkyl radical and Y is hydrogen or a functional group from the hydroxy, carboxy, amino, optionally low-alkyl- substituted mono or dialkylamino series, thiol, or a neg. acid group from the ester, phosphate, phosphonate, sulfate, sulfonate, maleimide, succinimidyl, epoxy, acrylate series. A biol., biochem. or synthetic indicator element can be coupled to B or Y as addn. or substitution reaction, whereby compds. can also be added imparting on the substrate surface a resistance against protein absorption and/or cell adhesion and in the B chain can be, optionally, composed of one or more ethylene oxide groups rather than one or more CH2 groups. According to the invention, said pptn. occurs on the surfaces of the substrates of pure or mixed oxides, nitrides or carbides of metals and semi-conductors. The invention is characterized in that the water-sol. salts composed of formula (IA) or (IB) are used to treat said surfaces, esp. the surfaces of sensor platforms, implants and medical accessory devices. The invention also relates to the use thereof as part of coated sensor platforms, implants and medical accessory devices in addn. to novel organophosphoric acids and organophosphonic acids themselves. The optionally substituted compds. of general formula (IA) and (IB), wherein the groups B and Y have the above mentioned designations i.e. optionally substituted alkyl, alkenyl, alkynyl, aryl, aralkyl, hetaryl or hetaryl, are equally called organophosphoric acids or phosphonic acids.

L21 ANSWER 2 OF 35 CAPLUS COPYRIGHT 2002 ACS

- 2002:90103 Document No. 136:149873 IL-17 molecules and uses thereof.

 Medlock, Eugene; Yeh, Richard; Silbiger, Scott M.; Elliot, Gary S.;

 Nguyen, Hung Q.; Jing, Shuqian (Amgen, Inc., USA). PCT Int. Appl. WO
 2002008285 A2 20020131, 242 pp. DESIGNATED STATES: W: AE, AG, AL, AM,

 AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK,

 DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE,

 KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX,

 MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ,

 UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW:

 AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR,

 IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English).

 CODEN: PIXXD2. APPLICATION: WO 2001-US19861 20010621. PRIORITY: US
 2000-PV213125 20000622; US 2001-PV266159 20010202; US 2001-810384
 20010316.
- AB Novel IL-17 like polypeptides and nucleic acid mols. encoding the same. The invention also provides vectors, host cells, selective binding agents, and methods for producing IL-17 like polypeptides. Also provided for are methods for the treatment, diagnosis, amelioration, or prevention of diseases with IL-17 like polypeptides, agonists, or antagonists thereof.
- L21 ANSWER 3 OF 35 CAPLUS COPYRIGHT 2002 ACS
- 2002:10538 Document No. 136:84704 Thymic stromal lymphopoietin receptor molecules and uses thereof. Pandey, Akhilesh; Ozaki, Katsutoshi; Baumann, Heinz; Levin, Steven D.; Farr, Andrew G.; Ziegler, Steven F.; Leonard, Warren J.; Lodish, Harvey F. (Whitehead Institute for Biomedical Research, USA). PCT Int. Appl. WO 2002000723 A2 20020103, 133 pp. DESIGNATED

- STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2001-US20730 20010628. PRIORITY: US 2000-PV214658 20000628.
- The present invention provides Thymic Stromal Lymphopoietin Receptor (TSLPR) polypeptides and nucleic acid mols. encoding the same. The invention also provides selective binding agents, vectors, host cells, and methods for producing TSLPR polypeptides. The invention further provides pharmaceutical compns. and methods for the diagnosis, treatment, amelioration, and/or prevention of diseases, disorders, and conditions assocd. with TSLPR polypeptides.
- L21 ANSWER 4 OF 35 CAPLUS COPYRIGHT 2002 ACS
 2002:409180 Document No. 137:1563 CD20/IgE-receptor like molecules and uses thereof. Welcher, Andrew A.; Calzone, Frank J. (USA). U.S. Pat. Appl. Publ. US 2002064823 A1 20020530, 46 pp., Cont.-in-part of U. S. Ser. No. 723,258. (English). CODEN: USXXCO. APPLICATION: US 2001-821821 20010329. PRIORITY: US 2000-PV193728 20000330; US 2000-723258 20001127.
- AB Novel CD20/IgE-receptor like polypeptides and nucleic acid mols. encoding the same. The invention also provides vectors, host cells, agonists and antagonists (including selective binding agents), and methods for producing CD20/IgE-receptor like polypeptides. Also provided for are methods for the treatment, diagnosis, amelioration, or prevention of diseases with CD20/IgE-receptor like polypeptides.
- L21 ANSWER 5 OF 35 CAPLUS COPYRIGHT 2002 ACS
- 2002:294172 Document No. 136:324075 IL-17 receptor-like polypeptides, polynucleotides and antibodies for identification of agonists and antagonists and for diagnosis/treatment of immune diseases. Jing, Shuqian (USA). U.S. Pat. Appl. Publ. US 20020045213 Al 20020418, 54 pp., Cont.-in-part of U.S. Ser. No. 724,460. (English). CODEN: USXXCO. APPLICATION: US 2001-809567 20010315. PRIORITY: US 2000-PV189816 20000316; US 2000-724460 20001128.
- Disclosed are novel IL-17 receptor like polypeptides and nucleic acid mols. encoding the same. The invention also provides vectors, host cells, antibodies, antisense oligonucleotides, agonists and antagonists (including selective binding agents), and methods for producing IL-17 receptor like polypeptides. Also provided are methods for the treatment, diagnosis, amelioration, or prevention of diseases assocd. With IL-17 receptor like polypeptides, e.g. immunol. diseases, autoimmune diseases, inflammation, transplant rejection, allergies, infections, obesity, anorexia, cachexia, neuronal diseases, lung diseases, skin diseases, kidney diseases, bone diseases, vascular diseases, cancer, etc. The invention further provides method for identifying antibody, small mol., protein, peptide, lipid, carbohydrate that mimicking or antagonizing the biol. activity of IL-17 receptor-like mol.
- L21 ANSWER 6 OF 35 CAPLUS COPYRIGHT 2002 ACS
- 2002:241284 Document No. 136:261833 Sequence homologs of interleukin 17 and their use in diagnosis and treatment of immunol. diseases, inflammations and infections. Medlock, Eugene; Yeh, Richard; Silbiger, Scott M.; Elliott, Gary S.; Nguyen, Hung Q.; Jing, Shuqian (USA). U.S. Pat. Appl. Publ. US 20020037524 A1 20020328, 91 pp., Cont.-in-part of U.S. Ser. No. 810,384. (English). CODEN: USXXCO. APPLICATION: US 2001-886404 20010621. PRIORITY: US 2000-PV213125 20000622; US 2001-PV266159 20010202; US 2001-810384 20010316.
- AB Novel sequence homologs of IL-17 polypeptides (IL-17E) and nucleic acid mols. encoding the same are disclosed. The invention also provides vectors, host cells, antibodies and other selective binding agents, and

methods for producing IL-17 like polypeptides. Also provided for are methods for the treatment, diagnosis, amelioration, or prevention of diseases with IL-17 like polypeptides, agonists, or antagonists thereof. Methods of high throughput drug screening for effectors of IL-17 polypeptides are another embodiment of the present invention.

- L21 ANSWER 7 OF 35 CAPLUS COPYRIGHT 2002 ACS
 2002:271976 Document No. 136:274360 Osteoprotegerin in treatment of
 osteoporosis and other bone diseases. Boyle, William J.; Lacey, David L.;
 Calzone, Frank J.; Chang, Ming-Shi (Amgen Inc., USA). U.S. US 6369027 B1
 20020409, 117 pp., Cont. of U.S. Ser. No. 577,788. (English). CODEN:
 USXXAM. APPLICATION: US 1996-706945 19960903. PRIORITY: US 1995-577788
 19951222.
- The present invention discloses a novel secreted polypeptide, AΒ osteoprotegerin, which is a member of the tumor necrosis factor receptor superfamily and is involved in the regulation of bone metab. Also disclosed are rat, mouse and human nucleic acids encoding osteoprotegerin, polypeptides, recombinant vectors and host cells for expression, antibodies which bind OPG, and pharmaceutical compns. Expression of rat OPG cDNA in transgenic mouse showed increase in bone d., particularly in femurs, pelvic bones and vertebrae. C-terminal truncations of osteoprotegerin are provided that inhibit bone resorption. Specifically, amino acid residues 22-185 which comprise four cysteine-rich domains are required for osteoprotegerin activity. Furthermore, osteoprotegerin monomers may be linked by disulfide linkages and the dimeric form of OPG appears to predominate in transgenic mice, although trimeric forms may also exist. The polypeptides are used to treat bone diseases characterized by increased resorption such as osteoporosis.
- L21 ANSWER 8 OF 35 CAPLUS COPYRIGHT 2002 ACS
 2001:886197 Document No. 136:32779 Novel cysteine-knot growth factor
 superfamily member: Cloaked-2 protein from human and mouse, their
 recombinant production and use in therapeutics. Paszty, Christopher J.;
 Gao, Yongming (Amgen, Inc., USA). PCT Int. Appl. WO 2001092308 A2
 20011206, 171 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA,
 BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES,
 FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ,
 LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL,
 PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ,
 VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ,
 CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC,
 ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2.
 APPLICATION: WO 2001-US17478 20010529. PRIORITY: US 2000-PV208550
 20000601; US 2000-PV223542 20000804.
- The present invention provides protein and cDNA sequences for novel Cloaked-2 proteins from human and mouse, which belong to cysteine-knot growth factor superfamily with conserved cysteine-knot motifs (CxGxC or CxC). The mRNA tissue expression profile of human Cloaked-2 protein is provided. The invention also provides vectors, host cells, selective binding agents, and methods for producing Cloaked-2 polypeptides. Also provided for are methods for the treatment, diagnosis, amelioration, or prevention of diseases with Cloaked-2 polypeptides. The invention further provides antibodies specific to Cloaked-2 proteins useful in therapeutics.
- L21 ANSWER 9 OF 35 CAPLUS COPYRIGHT 2002 ACS
 2001:833368 Document No. 135:370651 Receptor from TNF family. Boyle,
 William J.; Hsu, Hailing (Amgen Inc., USA). PCT Int. Appl. WO 2001085782
 A2 20011115, 124 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ,
 BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES,
 FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ,
 LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL,
 PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN,
 YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF,

- CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2001-US4568 20010212. PRIORITY: US 2000-PV181800 20000211.
- A member of the tumor necrosis factor family and its receptor are ΑB described. This member is primarily expressed in B cells and its expression correlates to increases in the no. of B cells and Igs produced. The natural, preferred human ortholog is here called AGP-3R. is a type III transmembrane protein and has an amino terminal extracellular domain, a transmembrane domain, and a carboxy terminal intracellular domain. AGP-3R-related proteins of the invention may be membrane-assocd. or in sol. form, recombinantly produced or isolated after natural prodn. The invention provides for nucleic acids encoding such AGP-3R-related proteins, vectors and host cells expressing the polypeptides, and methods for producing recombinant proteins. Antibodies or fragments thereof that specifically bind the proteins are also provided. AGP-3R agonists and antagonists are useful for modulating B cell response and treating inflammatory, immunol., and autoimmune diseases, e.g. rheumatoid arthritis, graft vs. host disease, lupus erythematosus, and Crohn's disease.
- L21 ANSWER 10 OF 35 CAPLUS COPYRIGHT 2002 ACS
- 2001:747853 Document No. 135:302897 CD20/IgE-receptor like molecules and uses thereof. Welcher, Andrew A.; Calzone, Frank J. (Amgen, Inc., USA). PCT Int. Appl. WO 2001074903 A2 20011011, 145 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2001-US10048 20010329. PRIORITY: US 2000-PV193728 20000330; US 2000-723258 20001127.
- AB Novel CD20/IgE-receptor like polypeptides and nucleic acid mols. encoding the same. The invention also provides vectors, host cells, agonists and antagonists (including selective binding agents), and methods for producing CD20/IgE-receptor like polypeptides. Also provided for are methods for the treatment, diagnosis, amelioration, or prevention of diseases with CD20/IgE-receptor like polypeptides.
- L21 ANSWER 11 OF 35 CAPLUS COPYRIGHT 2002 ACS
- 2001:713560 Document No. 135:268321 Human and murine fibroblast growth factor receptor-like polypeptides and encoding nucleic acids. Saris, Christiaan M.; Mu, Sharon X.; Xia, Min; Boone, Thomas Charles; Covey, Todd (Amgen, Inc., USA). PCT Int. Appl. WO 2001070977 A2 20010927, 163 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2001-US9073 20010322. PRIORITY: US 2000-PV191379 20000322.
- AB The present invention provides fibroblast growth Factor receptor-like (FGFR-L) polypeptides and nucleic acid mols. encoding the same found in cDNA libraries from mouse and human tissues. The murine FIGR-L polypeptide comprises a 1587-bp open reading frame encoding a protein of 529 amino acids and possessing a potential signal peptide at its N-terminus. Tissue profiling data for expression of the FIGR-L gene are also provided based on Norther blot anal. and RNase protection assays. The invention also provides selective binding agents, vectors, host cells,

and methods for producing FGFR-L polypeptides. The invention further provides pharmaceutical compns. and methods for the diagnosis, treatment, amelioration, and/or prevention of diseases, disorders, and conditions assocd. with FGFR-L polypeptides.

- L21 ANSWER 12 OF 35 CAPLUS COPYRIGHT 2002 ACS
- 2001:693376 Document No. 135:271903 IL-17 receptor like molecules and uses thereof. Jing, Shuqian; Medlock, Eugene; Yeh, Richard; Silbiger, Scott M.; Elliot, Gary S.; Nguyen, Hung Q. (Amgen Inc., USA). PCT Int. Appl. WO 2001068705 A2 20010920, 239 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2001-US8688 20010316. PRIORITY: US 2000-PV189923 20000316; US 2000-PV204208 20000512; US 2000-723232 20001127; US 2001-PV266159 20010202.
- AB Novel IL-17 receptor like polypeptides and nucleic acid mols. encoding the same. The invention also provides vectors, host cells, agonists and antagonists (including selective binding agents), and methods for producing IL-17 receptor like polypeptides. Also provided for are methods for treatment, diagnosis, amelioration, or prevention of diseases assocd. with IL-17 receptor like polypeptides, e.g. immune system dysfunction, inflammation, cancer and infection.
- L21 ANSWER 13 OF 35 CAPLUS COPYRIGHT 2002 ACS
- 2001:661596 Document No. 135:237110 Cloning and characterization of
 chordin-like-2 protein genes from human and mouse, diagnostic and
 therapeutic use thereof. Zhang, Ke; Linh, Cam; Nakayama, Naoki (Amgen,
 Inc., USA). PCT Int. Appl. WO 2001064885 A1 20010907, 167 pp. DESIGNATED
 STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU,
 CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP,
 KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX,
 NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG,
 US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE,
 BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT,
 LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN:
 PIXXD2. APPLICATION: WO 2001-US6891 20010302. PRIORITY: US 2000-PV186462
 20000302.
- The invention provides protein and cDNA sequences for novel human and mouse chordin-like-2 protein CHL2, which has sequence similarity to chordin known as the bone morphogenetic protein (BMP) inhibitor. The invention also provides selective binding agents, vectors, host cells, and methods for producing CHL2 polyproteins. The tissue distribution pattern of the mRNA shows that CHL2 is involved in mouse articular chondrocytes, sternum, placenta, uterus, colon, and small intestine. CHL2 directly interacts with BMPs, and its inhibitory activity is demonstrated in CHL2 gene transfected cells, Xenopus embryo, and in transgenic mice. The murine CHL2 gene is mapped to chromosome 7 centromere. The invention further provides pharmaceutical compns. and methods for the diagnosis, treatment, amelioration, and/or prevention of diseases, disorders, and conditions assocd. with CHL2 polyproteins.
- L21 ANSWER 14 OF 35 CAPLUS COPYRIGHT 2002 ACS
- 2001:472523 Document No. 135:66255 Liquid composition of a biodegradable
 block copolymer for drug delivery system. Seo, Min-hyo; Choi, In-ja
 (Samyang Corp., S. Korea). PCT Int. Appl. WO 2001045742 A1 20010628, 37
 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR,
 BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE,
 GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KZ, LC, LK, LR, LS, LT,

LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2000-KR1508 20001221. PRIORITY: KR 1999-60349 19991222. The present invention relates to a liq. polymeric compn. capable of forming a physiol. active substance-contg. implant when it is injected into a living body and a method of prepn. The compn. comprises a water-sol. biocompatible liq. polyethylene glycol deriv., a biodegradable block copolymer which is insol. in water but sol. in the water-sol. biocompatible liq. polyethylene glycol deriv. and a physiol. active substance. Thus, a triblock copolymer was prepd. from lactide-1,4-dioxanone and PEG. Piroxicam 150, the above biodegradable block copolymer 400, diacetyl polyethylene glycol 420, and gelatin 30 mg were dissolved in a 50% aq. HOAc soln. and the

drug-contg. liq. polymeric compn. was filtered and the org. solvent was

AΒ

removed.

L21 ANSWER 15 OF 35 CAPLUS COPYRIGHT 2002 ACS Document No. 134:331616 Sustained release microspheres based on 2001:300486 a carrier protein, a water soluble polymer and complexing agents. Scott, Terrence L.; Brown, Larry R.; Riske, Frank J.; Blizzard, Charles D.; Rashba-Step, Julia (Epic Therapeutics, Inc., USA). PCT Int. Appl. WO 2001028524 A1 20010426, 71 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 2000-US28200 20001012. PRIORITY: US 1999-420361 19991018.

A microsphere compn. for sustained release of therapeutic or diagnostic AΒ agents comprises (1) a carrier protein, (2) a water-sol . polymer, (3) a polyanionic polysaccharide as a first complexing agent, and (4) a divalent metal cation (Ca and Mg) as a second complexing agent. The microspheres have a smooth surface that includes a plurality of channel openings that are < 1000 .ANG. in diam. Various drugs were encapsulated into microspheres. For example, microspheres contg. leuprolide acetate were prepd. using human serum albumin (HSA), dextran sulfate, polyethylene glycol, and polyvinylpyrrolidone. The microspheres were composed of approx. 10% leuprolide acetate, 50% human serum albumin, 20% dextran sulfate and 20% polyethylene glycol/polyvinylpyrrolidone. Similar particles were prepd. which also included zinc sulfate or caprylic acid, both of which retarded the release of protein and peptide from the microspheres. Also, rifampicin-contg. HSA microspheres were prepd. with HSA incorporation of 74% and rifampicin incorporation into the particles of > 6.8%. The av. size of the particles was detd. to be 68 nm in diam.

L21 ANSWER 16 OF 35 CAPLUS COPYRIGHT 2002 ACS
2001:208500 Document No. 134:219363 Method for preparing electrophoresis supporter gel. Hayashizaki, Yoshihide (Riken Corp., Japan). PCT Int. Appl. WO 2001020317 A1 20010322, 30 pp. DESIGNATED STATES: W: CA, JP, US; RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE. (Japanese). CODEN: PIXXD2. APPLICATION: WO 2000-JP6247 20000913. PRIORITY: JP 1999-259013 19990913.

AB A method is described for prepg. an electrophoresis supporter (e.g., gel, entangled polymer) by cleaning using a weak alk. soln. (e.g.,

carbonate) at least a part of the surface of a supporting member (e.g., capillary column) contg. silicon used for supporting the electrophoresis supporter, and then, rendering the supporting member to carry the supporter. An electrophoresis gel is composed of a polyacrylamide-type polymer obtained by polymg. acrylamide or its deriv. (e.g., N,N-dimethylacrylamide, N-(hydroxymethyl)acrylamide) in the presence of two or more than two polar org. solvents (e.g., formamide, methanol). A method for efficiently sepg. nucleic acid (e.g., DNA, RNA) or PNA fragments by electrophoresis using the gel prepd. by the above method is claimed.

L21 ANSWER 17 OF 35 CAPLUS COPYRIGHT 2002 ACS

2001:168055 Document No. 134:208364 Amphipathic compound having dendritic structure. Tsuchida, Eishun; Takeoka, Shinji; Sou, Keitaro; Ohkawa, Haruki (Japan Science and Technology Corporation, Japan). PCT Int. Appl. WO 2001016211 A1 20010308, 60 pp. DESIGNATED STATES: W: US; RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE. (Japanese). CODEN: PIXXD2. APPLICATION: WO 2000-JP5702 20000824. PRIORITY: JP 1999-245731 19990831.

GI

$$R_0$$
 R_1 R_2 R_2

An amphipathic compd. having a dendritic structure represented by structural formula (I). In the I, R0 is a hydrophilic group (e.g., oligosaccharides); R1 and R2 each independently is a hydrophobic group; and n is an integer of 1 to 4. This amphipathic compd. can take advantage of the intermol. interaction to stably fix a water-sol.

polymer on the surface and can hold the same while retaining its intact function. Thus, a low generation dendritic compd. was prepd. by using lysine as a spacer, polyethylene oxide as the hydrophilic moiety former, and palmitic acid as the hydrophobic moiety former.

L21 ANSWER 18 OF 35 CAPLUS COPYRIGHT 2002 ACS

2001:468203 Document No. 135:66201 Conjugates targeted to the interleukin-2 receptor. Prakash, Ramesh K.; Clemens, Christopher M. (Watson Laboratories, Inc., USA). U.S. US 6251866 B1 20010626, 22 pp., Cont.-in-part of U.S. Ser. No. 914,042, abandoned. (English). CODEN: USXXAM. APPLICATION: US 1998-128572 19980804. PRIORITY: US 1997-914042 19970805.

AB A compn. for intracellular delivery of a chem. agent into an interleukin-2-receptor-bearing cell, e.g. an activated T cell, includes a chem. agent and at least one copy of an interleukin-2-receptor-binding and endocytosis-inducing ligand coupled to a water sol.

polymer. The ligand binds to a receptor on the interleukin-2-receptor-bearing cell and elicits endocytosis of the compn. The compn. also preferably includes a spacer for coupling the chem. agent and the ligand to the polymer. Chem. agents can include cytotoxins, transforming nucleic acids, gene regulators, labels, antigens, drugs, and the like. A preferred water sol.

polymer is a polyalkylene oxide, such as polyethylene glycol and polyethylene oxide, and activated derivs. thereof. The compn. can further comprise a carrier such as another water sol. polymer, liposome, or particulate. Methods of

using these compns. for delivering a chem. agent in vivo or in vitro are also disclosed. A method of detecting a disease, such as T-cell lymphocytic leukemia, T-cell acute lymphoblastic leukemia, peripheral T-cell lymphoma, Hodgkin's disease, or non-Hodgkin's lymphoma, assocd. with elevated levels of sol. IL-2 receptor is also disclosed.

- L21 ANSWER 19 OF 35 CAPLUS COPYRIGHT 2002 ACS
- 2001:891554 Document No. 136:24952 Surfactant-free emulsion compositions manufactured with water-soluble polymers.

 Nakano, Mitsuru (Nihon B.E.E. K. K., Japan). Jpn. Kokai Tokkyo Koho JP 2001342108 A2 20011211, 8 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 2000-166727 20000602.
- AB The compns., useful for cosmetics, foods, and pharmaceuticals, contain water-sol. polymers and are manufd. by directing a jet of fluid along a 1st path and redirecting the fluid in a controlled flow along a new path using a structure placed in the 1st path, wherein the directions of the 1st and the new path are designated to cause shear force and cavitation in the fluid. Liq. paraffin 10.0, cetyl 2-ethylhexanoate 10.0, beeswax 2.5, spermaceti 2.5, Me polysiloxane 0.5, stearic acid 0.5, glycerin 10.0, Me p-hydroxybenzoate 0.2, gum arabic 0.5, H2O 63.29, and lecithin 0.01 wt.% were preliminary emulsified with a homogenizer and emulsified with DeBEE (emulsifying app.) to give an emulsion with av. particle size 343 nm.
- L21 ANSWER 20 OF 35 CAPLUS COPYRIGHT 2002 ACS
- 2001:579316 Document No. 135:134281 Reagent for measuring glycosylated protein. Komiyama, Kishisato (Wako Pure Chemical Industries, Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 2001215229 A2 20010810, 7 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 2000-24916 20000202.
- AB A method and a reagent are provided for enzymically measuring a particular glycosylated protein (e.g., glycosylated albumin, glycosylated globulin, glycosylated transferrin, glycosylated HDL, glycosylated LDL, glycosylated anti-trypsin) in a biol. sample (e.g., whole blood, blood serum, blood plasma, urine). The particular glycosylated protein is specifically detd. based on the quantity of glycosylated amino acid or/and glycosylated peptide produced upon contacting the sample with a proteinase in the presence of a co-existing substance (e.g., antibodies, water -sol. polymer) capable of inhibiting the involvement of glycosylated proteins other than the particular glycosylated protein in the reaction with the proteinase. This method is superior to the conventional HPLC method in its versatility and rapidness.
- L21 ANSWER 21 OF 35 CAPLUS COPYRIGHT 2002 ACS
- 2001:729702 Document No. 135:278032 Polymer-based solid oral dosage forms with sustained drug release and high mechanical stability. Kolter, Karl; Schoenherr, Michael; Ascherl, Hermann (Basf A.-G., Germany). Eur. Pat. Appl. EP 1138321 A2 20011004, 14 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO. (German). CODEN: EPXXDW. APPLICATION: EP 2001-105547 20010306. PRIORITY: DE 2000-10015479 20000329.
- AB Solid oral dosage forms with sustained release characteristics comprise a drug, a mixt. of poly(vinyl acetate) and PVP, water-sol.

 polymers, and /or low- or high-mol. wt. lipophilic additives.
 Thus, tablets were prepd. from caffeine 160, Kollidon SR 160, Kollidon VA64 80 and Mg stearate 1.8 mg. The friability of tablets was <0.01% and the breaking strength was >325N.
- L21 ANSWER 22 OF 35 CAPLUS COPYRIGHT 2002 ACS
- 2001:923219 Document No. 136:42852 Preparation of oral sustained-release solid drug dosage forms. Kolter, Karl; Flick, Dieter; Ascherl, Hermann (BASF A.-G., Germany). Ger. Offen. DE 10029201 A1 20011220, 14 pp. (German). CODEN: GWXXBX. APPLICATION: DE 2000-10029201 20000619.
- AB Solid oral dosage forms with sustained release properties, contain at

least 1 drug, a preformulated mixt. from poly(vinyl acetate) and polyvinylpyrrolidone, optionally water-sol. polymers or lipophilic additives as well as the usual excipients. Granules obtained from the above mixt. are tabletted. Thus, a compn. contg. 400 g Kollidone SR/paracetamol mixt. (1:1) was granulated and the granules were mixed with 0.5% Mg stearate and compressed to give tablets.

- L21 ANSWER 23 OF 35 CAPLUS COPYRIGHT 2002 ACS
 2000:116860 Document No. 132:171073 Conjugates targeted to target receptors and/or interleukin-2 receptors. Prakash, Ramesh K.; Clemens, Christopher M. (Watson Laboratories, Inc.-Utah, USA). PCT Int. Appl. WO 2000007543 A2 20000217, 67 pp. DESIGNATED STATES: W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1999-US17648 19990804. PRIORITY: US 1998-128572 19980804.
- A compn. for intracellular delivery of a chem. agent into a target receptor and/or interleukin-2-receptor-bearing cell, e.g. an activated T cell and cancer cell, includes a chem. agent, at least one copy of target-receptor binding and/or an interleukin-2-receptor-binding and endocytosis-inducing ligand coupled to a water sol. polymer. The ligand binds to a target receptor and/or IL-2 receptor on the target receptor and/or IL-2-receptor-bearing cell and elicits endocytosis of the compn. The compn. also optionally includes a biodegradable spacer for coupling the chem. agent and the ligand to the polymer. Chem. agents can include cytotoxins, transforming nucleic acids, gene regulators, labels, antigens, drugs, and the like. A preferred water sol. polymer is polyalkylene oxide, such as polyethylene glycol and polyethylene oxide, and activated derivs. thereof. The compn. can further comprise a carrier such as another water sol. polymer, liposome, or particulate. Methods of using these compns. for delivering a chem. agent in vivo or in vitro are also disclosed. A method of detecting a disease, such as cancer, T-cell lymphocytic leukemia, T-cell acute lymphoblastic leukemia, peripheral T-cell lymphoma, Hodgkin's disease, and non-Hodgkin's lymphoma, assocd. with elevated levels of sol. target receptor and/or IL-2 receptor is also disclosed.
- L21 ANSWER 24 OF 35 CAPLUS COPYRIGHT 2002 ACS Document No. 132:177439 Method for production of physiologically 2000:139183 active peptides. Hirata, Akira; Murakami, Yoshihiko (Kobe Steel, Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 2000063399 A2 20000229, 6 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1998-236055 19980821. The title method comprises (1) adding the protease and the protein to a AB 2-phase water-sol. polymer system to hydrolyze the protein; (2) removing the physiol. active peptide (i.e., the product) from one phase of the 2-phase watersol. polymer system. The 2-phase polymer system comprises polyethylene glycol and dextran. The enzymic hydrolysis of the protein is done in the polyethylene glycol rich phase, and the product is removed from the dextran rich phase. The title method is highly efficient.
- L21 ANSWER 25 OF 35 SCISEARCH COPYRIGHT 2002 ISI (R)
 2000:421268 The Genuine Article (R) Number: 319KX. The size of membrane
 pores: The effect of non-electrolytes on the conductance of gramicidin.
 Coates G M P (Reprint); Alder G M; Smart O S; Bashford C L. UNIV
 BIRMINGHAM, SCH BIOSCI, BIRMINGHAM B15 2TT, W MIDLANDS, ENGLAND (Reprint);

ST GEORGE HOSP, SCH MED, DEPT BIOCHEM & IMMUNOL, LONDON SW17 ORE, ENGLAND. ACTA PHYSICA POLONICA B (MAY 2000) Vol. 31, No. 5, pp. 1097-1107. Publisher: ACTA PHYSICA POLONICA B, JAGELLONIAN UNIV, INST PHYSICS. REYMONTA 4, 30-059 KRAKOW, POLAND. ISSN: 0587-4254. Pub. country: ENGLAND. Language: English.

ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS

The effects of neutral polymers on ion channel conductance have been used in the past to estimate channel radius. We have measured the effect of Polyethylene-glycol and dextrans on gramicidin-D, a peptide ion channel. The availability of high resolution structures of gramicidin-ii allows us to make a direct comparison between the characteristic radius obtained by these experiments and the radius of the channel obtained from the NMR structure. The effects of PEG on gramicidin are significantly different from those observed on other, wider channels, and the experiment suggests that the operational size of the gramicidin channel exceeds that seen in the NMR and crystal structures. Our data using non-dehydrating polymers such as dextrans, provide estimates of gramicidin channel size smaller than those obtained with PEGs and closer to those predicted by the NMR and crystal structures.

L21 ANSWER 26 OF 35 CAPLUS COPYRIGHT 2002 ACS

1999:133618 Document No. 130:187175 Conjugates targeted to the interleukin-2 receptor. Prakash, Ramesh K. (Theratech, Inc., USA). PCT Int. Appl. WO 9907324 A2 19990218, 53 pp. DESIGNATED STATES: W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 1998-US16290 19980805. PRIORITY: US 1997-914042 19970805.

A compn. for intracellular delivery of a chem. agent into an AΒ interleukin-2-receptor-bearing cell, e.g. an activated T cell, includes a chem. agent and at least two copies of an interleukin-2-receptor-binding and endocytosis-inducing ligand coupled to a water sol . polymer. The ligand binds to a receptor on the interleukin-2-receptor-bearing cell and elicits endocytosis of the compn. The compn. also optionally includes a spacer for coupling the chem. agent and the ligand to the polymer. Chem. agents can include cytotoxins, transforming nucleic acids, gene regulators, labels, antigens, drugs, and the like. A preferred water sol. polymer is polyalkylene oxide, such as polyethylene glycol and polyethylene oxide, and activated derivs. thereof. compn. can further comprise a carrier such as another water sol. polymer, liposome, or particulate. Methods of using these compns. for delivering a chem. agent in vivo or in vitro are

L21 ANSWER 27 OF 35 CAPLUS COPYRIGHT 2002 ACS

also disclosed.

1999:751514 Document No. 131:356114 Temperature-responsive biodegradable polymers providing novel drug delivery systems. Yui, Nobuhiko (Foundation for Scientific Technology Promotion, Japan). Jpn. Kokai Tokkyo Koho JP 11322941 A2 19991126 Heisei, 7 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1998-127979 19980511.

The polymers, which are degraded in cells, tissues, or organs by enzymes and temp. increase and are useful as drug carriers, comprise (a) 3-dimensional net work structure of water-sol. biodegradable polymers or polymers having biodegradable site and (b) temp.-responsive polymers grafted on (a). Three N-isopropylacrylamide-N,N-dimethylacrylamide copolymers having amino group at one end (prepn. given, Mn 2600, 4200, or 8800) were treated

with methacryl chloride to give copolymers having methacryl group at the other terminal. A compn. contg. each copolymer, dextran methacrylate, ammonium persulfate, and DMSO was irradiated with UV at room temp. for 4 h to give a hydrogel, which was soaked in H2O at room temp. for 10 days. Transmittance of the swollen hydrogel in a phosphate buffer was decreased with increase in the temp. for all 3 hydrogels. Degrdn. of the swollen gel with dextranase was promoted with increase in the temp. for the hydrogel having grafted chain with Mn 4200 or 8800.

- L21 ANSWER 28 OF 35 CAPLUS COPYRIGHT 2002 ACS
- 1999:746965 Document No. 131:342078 Polyrotaxane supramolecular materials for implants. Yui, Nobuhiko; Ohtani, Akira (Foundation for Scientific Technology Promotion, Japan). Jpn. Kokai Tokkyo Koho JP 11319069 A2 19991124 Heisei, 5 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1998-127978 19980511.
- AB The materials comprise cyclic mols. threaded with biodegradable or hydrolyzable groups-terminated water-sol. linear macromols., where the cyclic mols. are not eliminated unless the terminal groups are decompd. Polyethylene glycol was esterified with succinic anhydride, imidized with N-hydroxysuccinimide, amidated with ethylenediamine, treated with .alpha.-cyclodextrin, and the resulting pseudopolyrotaxane was amidated with Z-Phe-OSu (Su = succinimido) to give a polyrotaxane having degrdn. point 298.0.degree.
- L21 ANSWER 29 OF 35 SCISEARCH COPYRIGHT 2002 ISI (R)
 1999:716175 The Genuine Article (R) Number: 236EY. Bioconjugation in
 pharmaceutical chemistry. Veronese F M (Reprint); Morpurgo M. UNIV PADUA,
 DEPT PHARMACEUT SCI, VIA F MARZOLO 5, I-35131 PADUA, ITALY (Reprint).
 FARMACO (30 AUG 1999) Vol. 54, No. 8, pp. 497-516. Publisher: ELSEVIER
 SCIENCE SA. PO BOX 564, 1001 LAUSANNE, SWITZERLAND. ISSN: 0014-827X. Pub.
 country: ITALY. Language: English.

ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS

AΒ

Polymer conjugation is of increasing interest in pharmaceutical chemistry for delivering drugs of simple structure or complex compounds such peptides, enzymes and oligonucleotides. For long time drugs, mainly with antitumoral activity, have been coupled to natural or synthetic polymers with the purpose of increasing their blood permanence time, taking advantage of the increased mass that reduces kidney ultrafiltration. However only recently complex constructs were devised that exploit the 'enhanced permeability and retention' (EPR) effect for an efficient tumor targeting, the high molecular weight for adsorption or receptor mediated endocytosis and finally a lysosomotropic targeting, taking advantage of acid labile bonds or cathepsin susceptible polypeptide spacers between polymer and drug. New original, very active conjugates of this type, as those based on poly(hydroxyacrylate) polymers, are already in advanced state of development. Labile oligonucleotides, including antisense drugs, were also successfully coupled to polymers in view of an increased cell penetration and stabilization towards nucleases. However, the most active research activity resides in the field of polypeptides and proteins delivery, mainly for the two following reasons: first of all because a great number of therapeutically interesting compounds are now being produced by genetic engineering in large quantity and, secondly, because these products are difficult to administer to patients for several inherent drawbacks. Proteins are in fact easily digested by many endo- and exo-peptidases present in blood or in other body districts; most of them are immunogenic to some extent and, finally, they are rapidly excreted by kidney ultrafiltration. Covalent polymer conjugation at protein surface was demonstrated to reduce or eliminate these problems, since the bound polymer behaves like a shield hindering the approach of proteolytic enzymes, antibodies, or antigen processing cell. Furthermore, the increase of the molecular weight of the conjugate allows to overcome the kidney elimination threshold. Many successful results were already

obtained in peptides and proteins, conjugated mainly to water soluble or amphiphilic polymers like poly(ethylene glycol) (PEG), dextrans, or styrene-maleic acid anhydride. Among the most successful are the conjugates of asparaginase, interleukin-2 or -6 and neocarcinostatin, to remind some antitumor agents, adenosine deaminase employed in a genetic desease treatment, superoxide dismutase as scavenger of toxic radicals, hemoglobin as oxygen carrier and urokinase and streptokinase as proteins with antithrombotic activity. In pharmaceutical chemistry the conjugation with polymers is also of great importance for synthetic applications since many enzymes without loss of catalytic activity become soluble in organic solvents where many drug precursors are. The various and often difficult chemical problems encountered in conjugation of so many different products prompted the development of many synthetic procedures, all characterized by high specificity and mild condition of reaction, now known as 'bioconjugation chemistry'. Bioconjugation developed also the design of new tailor-made polymers with the wanted molecular weight, shape, structure and with the functional groups needed for coupling at the wanted positions in the chain. (C) 1999 Elsevier Science S.A. All rights reserved.

- L21 ANSWER 30 OF 35 MEDLINE
- 1999259607 Document Number: 99259607. PubMed ID: 10327623. Aqueous two-phase systems containing self-associating block copolymers. Partitioning of hydrophilic and hydrophobic biomolecules. Svensson M; Berggren K; Veide A; Tjerneld F. (Department of Physical Chemistry 1, Lund University, Sweden.) JOURNAL OF CHROMATOGRAPHY. A, (1999 Apr 16) 839 (1-2) 71-83. Journal code: 9318488. Pub. country: Netherlands. Language: English.
- A series of proteins and one membrane-bound peptide have been AΒ partitioned in aqueous two-phase systems consisting of micelle-forming block copolymers from the family of Pluronic block copolymers as one polymer component and dextran T500 as the other component. The Pluronic molecule is a triblock copolymer of the type PEO-PPO-PEO, where PEO and PPO are poly(ethylene oxide) and poly(propylene oxide), respectively. Two different Pluronic copolymers were used, P105 and F68, and the phase diagrams were determined at 30 degrees C for these polymer systems. Since the temperature is an important parameter in Pluronic systems (the block copolymers form micellar-like aggregates at higher temperatures) the partitioning experiments were performed at 5 and 30 degrees C, to explore the effect of temperature-triggered micellization on the partitioning behaviour. The temperatures correspond to the unimeric (single Pluronic chain) and the micellar states of the P105 polymer at the concentrations used. The degree of micellization in the F68 system was lower than that in the P105 system, as revealed by the phase behaviour. A membrane-bound peptide, gramicidin D, and five different proteins were partitioned in the above systems. The proteins were lysozyme, bovine serum albumin, cytochrome c, bacteriorhodopsin and the engineered B domain of staphylococcal protein A, named Z. The Z domain was modified with tryptophan-rich peptide chains in the C-terminal end. It was found that effects of salt dominated over the temperature effect for the water-soluble proteins lysozyme, bovine serum albumin and cytochrome c. A strong temperature effect was observed in the partitioning of the integral membrane protein bacteriorhodopsin, where partitioning towards the more hydrophobic Pluronic phase was higher at 30 degrees C than at 5 degrees C. The membrane-bound peptide gramicidin D partitioned exclusively to the Pluronic phase at both temperatures. The following trends were observed in the partitioning of the Z protein. (i) At the higher temperature, insertion of tryptophan-rich peptides increased the partitioning to the Pluronic phase. (ii) At the lower temperature, lower values of K were observed for ZT2 than for ZT1.

- 1998:761809 Document No. 130:17218 Targeted delivery to T lymphocytes.
 Prakash, Ramesh K.; Kumar, Vijay (Theratech, Inc., USA). PCT Int. Appl.
 WO 9851336 A1 19981119, 46 pp. DESIGNATED STATES: W: AL, AM, AT, AU, AZ,
 BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH,
 GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU,
 LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK,
 SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU,
 TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR,
 GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG.
 (English). CODEN: PIXXD2. APPLICATION: WO 1998-US9057 19980504.
 PRIORITY: US 1997-857009 19970515.
- AB A compn. for intracellular delivery of a chem. agent into a T cell comprises a receptor-binding and endocytosis-inducing ligand and a chem. agent coupled to a water sol. polymer. The ligand binds to a receptor on T lymphocytes and elicits endocytosis of the compn. The compn. also includes a spacer for coupling the chem. agent and the ligand to the polymer. Chem. agents can include cytotoxins, transforming nucleic acids, gene regulators, labels, antigens, drugs, and the like. A preferred water sol. polymer is polyethyleneglycol and activated derivs. thereof. The compn. can further comprise a carrier such as a water sol. polymer, liposome, or particulate. Methods of using these compns. for delivering a chem. agent in vivo or in vitro are also disclosed.
- L21 ANSWER 32 OF 35 CAPLUS COPYRIGHT 2002 ACS
 1996:676099 Document No. 125:309046 Drug release systems containing
 water-soluble polymer domain and biodegradable
 hydrogel as matrix. Yui, Nobuhiko (Shingijutsu Kaihatsu Jigyodan, Japan).

 Jpn. Kokai Tokkyo Koho JP 08231435 A2 19960910 Heisei, 5 pp.
 (Japanese). CODEN: JKXXAF. APPLICATION: JP 1995-38427 19950227.
- AB Stimulation-responsive drug release systems comprise watersol. polymer domain (e.g. polyethylene
 glycol) and biodegradable hydrogel (e.g. dextran) as
 matrix. Active ingredients such as insulin showed selective distribution
 in the polyethylene glycol-dextran diphase.
 Active ingredients (e.g. insulin) as well as the polymer domain
 are released in response to biodegradable hydrogel from the
 surface.
- L21 ANSWER 33 OF 35 CAPLUS COPYRIGHT 2002 ACS
 1995:319762 Document No. 122:89553 PEG hydrazone and PEG oxime linkage
 forming reagents and protein derivatives. Wright, David E. (Ortho
 Pharmaceutical Corp., USA). Eur. Pat. Appl. EP 605963 A2 19940713, 47 pp.
 DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU,
 MC, NL, PT, SE. (English). CODEN: EPXXDW. APPLICATION: EP 1993-309825
 19931207. PRIORITY: US 1992-987739 19921209; US 1993-45052 19930407; US
 1993-157343 19931123.
- Compds. for modifying polypeptides with PEG or other watersol. org. polymers are described. The watersol. polymer reagents include hydrazine, hydrazine
 carboxylate, semicarbazole, thiosemicarbazide, carbonic acid dihydrazide,
 carbazide, thiocarbazide, and arylhydrazide derivs. as well as oxylamine
 derivs. of water-sol. org. polymers, such as
 polyethylene glycol, polypropylene glycol,
 polyoxyethylated polyol, heparin, heparin fragments, dextran
 polysaccharides, polyamino acids, and polyvinyl alc. Kits for modifying
 polypeptides with the above water-sol. polymer
 reagents are also provided. Thus, erythropoietin was modified by oxidn.
 and treatment with monomethoxypolyoxyethylene semicarbazide and the
 product was sepd. by chromatog. The antigenicity and the effect on
 hematocrit levels of the above derivs. were demonstrated.

- 1992:190638 Document No. 116:190638 Separation of mixtures by two-phase systems. Hsu, James T. (Lehigh University, USA). U.S. US 5078886 A 19920107, 10 pp. Cont.-in-part of U.S. 4,980,065. (English). CODEN: USXXAM. APPLICATION: US 1990-591832 19901002. PRIORITY: US 1989-423333 19891018.
- An org. 2-phase system is useful for the sepn. and purifn. of chems., biochems., and optical isomers. The 2-phase system can be formed with water-sol. polymers as 1 phase, and a chiral compd. as the other phase together with a water-miscible org. solvent and/or water. A L-lysine/PEG 8000/H2O system was prepd. and used to sep. D- and L-phenylalanine. L-Phenylalanine gave a lower partition coeff. than D-phenylalanine and was therefore enriched in the L-lysine (lower) phase.
- L21 ANSWER 35 OF 35 CAPLUS COPYRIGHT 2002 ACS
 1991:488820 Document No. 115:88820 Two-phase system comprising chiral compound and water soluble polymer for mixture separation. Hsu, James T. (Lehigh University, USA). U.S. US 4980065 A 19901225, 8 pp. (English). CODEN: USXXAM. APPLICATION: US 1989-423333 19891018.
- An aq. two-phase system consisting of (1) a chiral compd., (2) a AΒ water sol. polymer, and (3) water is prepd. for sepg. optical isomer or stereoisomer of (in)org. compds. or biomaterial, e.g. protein, peptide, cells and cell particles. The chiral compd. is a D- or L-.alpha.-amino acid or monosaccharide, or disaccharide or chiral salt, or chiral water miscible solvent or chiral acid or chiral base. The water sol. polymer is a polypropylene glycol (mol. wt. 300-50,000), poly(vinylpyrrolidone), poly(vinylalc.), dextran, and sodium dextran sulfate. Thus, 28.57% wt./wt. L-lysine, 10.71% wt./wt. polyethylene glycol 8000, and 60.72% wt./wt. water were mixed to form a two-phase system for the sepn. of D- and L-phenylalanine, .beta.-lactoglobulins A and B, and D- and L-tryptophan with D-phenylalanine, .beta.-lactoglobulin A, and D-tryptophan in favor in upper phase. The invention provides an improved process for affinity partitioning and for partition affinity ligand assay. Method studies counter-current distribution, cross-current extn., or counter-current extn. are also included in the process for anal., preparative, and large scale com. sepn. using the two-phase system.

=> s modified Apo A I L23 12 MODIFIED APO A I

=> dup remove 123
PROCESSING COMPLETED FOR L23
L24 4 DUP REMOVE L23 (8 DUPLICATES REMOVED)

=> d 124 1-4 cbib abs

L24 ANSWER 1 OF 4 MEDLINE DUPLICATE 1
97382910 Document Number: 97382910. PubMed ID: 9240901. Optimized automated apolipoprotein A-I assays as markers for coronary artery disease. Levinson S S; Hobbs G A. (Department of Veterans Affairs Medical Center, and the Department of Pathology, University of Louisville, KY 40206, USA.) ARCHIVES OF PATHOLOGY AND LABORATORY MEDICINE, (1997 Jul) 121 (7) 678-84. Journal code: 7607091. ISSN: 0003-9985. Pub. country: United States. Language: English.

AB BACKGROUND: Studies are divided as to whether or not apolipoprotein A-I (apo A-I) is a better marker for coronary artery disease (CAD) than

high-density lipoprotein cholesterol. We hypothesized that the detergent Tween 20, which is thought to expose antigenic sites in apo A-I, would improve automated kit apo A-I assays as a diagnostic marker for CAD. METHODS: Apolipoprotein A-I was assayed by two standard automated methods and by the same methods after serum samples and reagents had been treated with Tween 20. Serum samples were obtained from 226 consecutive male patients, age 40-70 years, presenting for angiography, except for defined exclusion characteristics. Patients were categorized into two groups on the basis of stenosis: (1) normal, all vessels <20% stenosis, $\tilde{n}=79$, and (2) CAD, at least one vessel >70% stenosis, n = 147. Diagnostic accuracy was assessed by receiver operator characteristic stenosis curves and forward stepwise logistic regression, where adjustment was made for significant possible confounding characteristics and drugs. RESULTS: The optimal concentration of Tween 20 was found to be 0.5%. Receiver operator characteristic curves showed a greater area for apo A-I with Tween (area = 0.63 to 0.64) as compared to apo A-I without Tween (area = 0.60 to 0.62). Logistic regression indicated that apo A-I with Tween was a significantly better marker than high-density lipoprotein cholesterol. Receiver operator characteristic curves indicated that the ratio of modified

apo A-I to apo B gave a significant
improvement in area over the ratio of high-density to low-density
lipoprotein cholesterol. CONCLUSIONS: Addition of Tween 20 to apo A-I
assays improved diagnostic discrimination for CAD. The modified

apo A-I assays were better markers than high-density lipoprotein cholesterol, and the ratio of apolipoproteins was significantly better markers than lipoprotein lipids. These findings may explain the discrepancies between studies comparing high-density lipoprotein cholesterol and apo A-I as markers for CAD. Our data suggest that a multicenter effort toward optimizing and clinically validating apo A-I test reagents may be worthwhile.

L24 ANSWER 2 OF 4 MEDLINE DUPLICATE 2
96206557 Document Number: 96206557. PubMed ID: 8621158. Binding of
apolipoprotein A-I and acetaldehyde-modified apolipoprotein A-I to liver
extracellular matrix. Paradis V; Mathurin P; Ratziu V; Poynard T; Bedossa
P. (Service d'Anatomie Pathologique, Hopital de Bicetre, Le
Kremlin-Bicetre, Paris, France.) HEPATOLOGY, (1996 May) 23 (5) 1232-8.
Journal code: 8302946. ISSN: 0270-9139. Pub. country: United States.
Language: English.

Apolipoprotein A-I (Apo A-I), a protein produced mainly by hepatocytes, is AΒ decreased in the sera of alcoholic patients with liver fibrosis and cirrhosis. To explain this decrease, we investigated possible interactions between liver extracellular matrix (ECM) and Apo A-I. Using a solid-phase binding assay, we evaluated the binding of Apo A-I to the different liver matrix components. Apo A-I bound significantly to fibronectin (FN) (optical density [OD] = 1.11 +/- .26, P = .01) and collagen (C) I (OD = 0.91 + -0.22, P = .02) in comparison with bovine serum albumin (BSA) (OD = 0.26 + /- 0.16). Binding of Apo A-I to fibronectin was concentration dependent and saturable. Apo A-I bound also to ECM in vivo because Apo A-I was detected by immunofluorescence on fibrous septa in liver biopsy specimens of alcoholic patients. Because a negative correlation between Apo A-I and liver fibrosis is amplified in alcoholic patients, we investigated whether the in vitro formation of Apo A-I/acetaldehyde complex (adducts) increased the binding of Apo A-I to the ECM. We showed that the amount of Apo A-I that bound to FN was significantly higher with acetaldehyde-modified Apo A-I (OD

= 2.18 + /- 0.19, P = .01) than with native Apo A-I. This increase was probably related to the formation and binding of Apo A-I dimers, because immunoblot of in vitro acetaldehyde-modified Apo

A-I showed the formation of dimeric Apo A-I. In conclusion, FN binds both native and acetaldehyde-modified

Apo A-I. Because FN is deposited early and in excess during liver fibrosis, a storage mechanism of Apo A-I on newly

deposited fibronectin would explain, in part, the decrease observed in alcoholic patients with liver fibrosis.

L24 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2002 ACS Document No. 122:299055 Cholesteryl ester transfer protein 1995:541414 inhibitor polypeptide, antibodies against the synthetic polypeptide and prophylactic and therapeutic anti-atherosclerosis treatments. Kushwaha, Rampratap S.; McGill, Henry C., Jr.; Kanda, Patrick (Southwest Foundation for Biomedical Research, USA). PCT Int. Appl. WO 9504755 A1 19950216, 47 pp. DESIGNATED STATES: W: AU, CA, JP, KR; RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE. (English). CODEN: PIXXD2. APPLICATION: WO 1994-US8624 19940802. PRIORITY: US 1993-102160 19930804. A polypeptide and analogs thereof inhibit cholesteryl ester transfer AΒ protein (CETP). An anti-atherosclerosis compn. comprises an anti-atherosclerosis effective amt. of the polypeptide and a pharmaceutically acceptable carrier. An anti-atherosclerosis kit comprises in sep. sterile containers at least one unit of the compn. contg. the polypeptide, one syringe, and one needle. An antibody has specificity for the polypeptide of the invention, the baboon CETP 4-kDa polypeptide inhibitor, the 1-36 amino acid N-terminal fragment of apoC-I, modified apo A-I (mol. wt. 31 kDa), or modified apoE (mol. wt. 41 kDa). A method of preventing atherosclerosis in a mammal being predisposed to that condition comprises administering to the mammal a prophylactically effective amt. of the polypeptide of the invention, and a method of treating a mammal afflicted with atherosclerosis comprises the administration of a therapeutically effective amt. of the polypeptide. The peptides consist of the N-terminal 36 residues of baboon apoC-I, a synthetic peptide adding an N-terminal Ala-Pro dipeptide, the human 38-residue analog, and active fragments and substituted analogs.

L24 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2002 ACS
1993:552092 Document No. 119:152092 Cholesteryl ester transfer protein
(CETP) inhibitor polypeptide, antibodies against the synthetic
polypeptide, and prophylactic and therapeutic anti-atherosclerosis
treatments. Kushwaha, Rampratap; Born, Kathleen; Mcgill, Henry C., Jr.;
Kanda, Patrick; Dunham, Raymond G. (Southwest Foundation for Biomedical
Research, USA). PCT Int. Appl. WO 9311782 A1 19930624, 46 pp. DESIGNATED
STATES: RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT,
SE. (English). CODEN: PIXXD2. APPLICATION: WO 1992-US10846 19921215.
PRIORITY: US 1991-811049 19911219.

AB A polypeptide and analogs thereof inhibit CETP. Also disclosed are methods using the polypeptide of the invention for prevention and treatment of atherosclerosis, an anti-atherosclerosis compn., and an anti-atherosclerosis kit. An antibody is disclosed which has specificity for the polypeptide of the invention, the baboon CETP 4 kDa polypeptide inhibitor, the 1-36 apo C-I amino-terminal fragment, modified apo A-I (31 kDa mol. wt.), or modified apo E (41 kDa mol. wt.). Sequences of polypeptide inhibitors are included. Detection of the CETP inhibitor peptide in the plasma of baboons with a high HDL1 phenotype is described, as is CETP inhibition by various peptide fragments.

=> s fusion protein L25 133094 FUSION PROTEIN

=> s 125 chimeric MISSING OPERATOR L25 CHIMERIC The search profile that was entered contains terms or nested terms that are not separated by a logical operator.

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17559 L25 AND CHIMERIC
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=> s 126 and IgG

L26

L27 723 L26 AND IGG

=> s 127 and Apo A I

L28 0 L27 AND APO A I

=> s 127 and human Apo-A-I

3 FILES SEARCHED...

L29 0 L27 AND HUMAN APO-A-I

=> s 127 and apolipoprotein AI

L30 0 L27 AND APOLIPOPROTEIN AI

=> s (dayer j?/au or burger d?/au or kohno t?/au or edwards c?/au)
L31 14667 (DAYER J?/AU OR BURGER D?/AU OR KOHNO T?/AU OR EDWARDS C?/AU)

=> s 131 and ApoAI

L32 1 L31 AND APOAI

=> d 132 cbib abs

AB

L32 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS
2001:798252 Document No. 135:362518 Apo-AI/AII peptide derivatives for hypocholesteremic and antiviral therapy. Kohno, Tadahiko (Amgen Inc., USA). PCT Int. Appl. WO 2001081376 A2 20011101, 49 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2001-US13068 20010423. PRIORITY: US 2000-PV198920 20000421.

The present invention concerns therapeutic agents that mimic the activity of Apo-AI amphipathic helix peptide. In accordance with the present invention, the compds. of the invention comprise: (a) a Apo-AI amphipathic helix peptide or Apo-AI amphipathic helix peptide-mimetic domain, preferably the amino acid sequence of SEQ ID NO:7, or sequences derived therefrom by phage display, RNA-peptide screening, or the other techniques mentioned above; and (b) a vehicle, such as a polymer (e.g., PEG or dextran) or an Fc domain, which is preferred; wherein the vehicle, preferably an Fc domain, is covalently attached to the Apo-AI amphipathic helix peptide or Apo-AI amphipathic helix peptide-mimetic domain. The vehicle and the Apo-AI amphipathic helix peptide or Apo-AI amphipathic helix peptide-mimetic domain may be linked through the N- or C-terminus of the Apo-AI amphipathic helix peptide or Apo-AI amphipathic helix peptide-mimetic domain, as described further below. The preferred vehicle is an Fc domain, and the preferred Fc domain is an IgG Fc domain. Preferred Apo-AI amphipathic helix peptide or Apo-AI amphipathic helix peptide-mimetic domains comprise the amino acid sequences described in Table 1. Other Apo-AI amphipathic helix peptide or Apo-AI amphipathic helix peptide-mimetic domains can be generated by phage display, RNA-peptide screening and the other techniques mentioned herein.

L33 0 L31 AND HUMAN APO-A-I

^{=&}gt; s 131 and human Apo-A-I 3 FILES SEARCHED...

AΒ

=> s l1 and lipoprotein L35 142 L1 AND LIPOPROTEIN

=> dup remove 135 PROCESSING COMPLETED FOR L35 L36 124 DUP REMOVE L35 (18 DUPLICATES REMOVED)

=> s 136 and modified 1.37 1 L36 AND MODIFIED

=> d 137 cbib abs

L37 ANSWER 1 OF 1 SCISEARCH COPYRIGHT 2002 ISI (R)
93:736313 The Genuine Article (R) Number: MM035. ENDOCYTOSIS AND DEGRADATION
OF BOVINE APO-LACTOFERRIN AND HOLO-LACTOFERRIN BY ISOLATED RAT HEPATOCYTES
ARE MEDIATED BY RECYCLING CALCIUM-DEPENDENT BINDING-SITES. MCABEE D D
(Reprint); NOWATZKE W; OEHLER C; SITARAM M; SBASCHNIG E; OPFERMAN J T;
CARR J; ESBENSEN K. UNIV NOTRE DAME, DEPT BIOL SCI, NOTRE DAME, IN, 46556
(Reprint). BIOCHEMISTRY (14 DEC 1993) Vol. 32, No. 49, pp. 13749-13760.
ISSN: 0006-2960. Pub. country: USA. Language: ENGLISH.
ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS

We characterized endocytosis of iron-saturated(holo) and iron-depleted (apo) I-125-labeled bovine lactoferrin (Lf) by isolated rat hepatocytes. Hepatocytes ingested both Lf forms-determined by EGTA/dextran sulfate removal of surface-bound Lf-at maximal endocytic rates of 1.85 and 1.52 fmol cell-1 min-1 for I-125-apo-Lf and I-125-holo-Lf, respectively. First-order endocytic rate constants (37-degrees-C) for I-125-apo-Lf and I-125-holo-Lf were 0.276 and 0.292 min-1, respectively. Regardless of Lf's iron content, hyperosmotic media (approximately 500 mmol/kg) inhibited Lf uptake by approximately 90%, indicating endocytosis of both Lf forms was primarily clathrin-dependent. Endocytosis of both Lf forms was not altered significantly in the presence of excess iron chelator desferrioxamine or rat holo-transferrin, or by cycloheximide treatment. Fluorescein isothiocyanate- and cyclohexanedionemodified Lf competed fully with native Lf for binding and endocytosis, indicating that, unlike human Lf, modification of lysine or arginine residues does not block the interaction of bovine Lf with cells. After binding Lf at 4-degrees-C, cells at 37-degrees-C internalized approximately 90% of Lf bound to Ca2+-dependent sites but not Lf bound to Ca2+-independent sites. Following uptake, hepatocytes released acid-soluble (degraded) products of I-125-Lf biphasically at 37-degrees-C, an initial rapid phase within the first 20 min-more pronounced with I-125-holo-Lf-followed by a sustained linear release of 298 and 355 molecule equiv cell-1 min-1 for I-125-apo-Lf and I-125-holo-Lf, respectively. At 4-degrees-C, both digitonin-permeabilized and intact cells bound approximately 1.1 x 10(6) I-125-Lf molecules to Ca2+-dependent sites per cell, indicating that hepatocytes do not contain a sizeable intracellular pool of these sites. Moreover, cells retained >70% of Ca2+-dependent sites on the surface during sustained Lf endocytosis. Thus, these Lf binding sites recycle during endocytosis at an estimated 4-5 min/circuit.

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